of the voting shares of First Colonial Bank, Hopewell, Virginia, which is the proposed successor by charter conversion to First Colonial Bank, FSB, Hopewell, Virginia.

C. Federal Reserve Bank of San Francisco (Maria Villanueva, Manager of Analytical Support, Consumer Regulation Group) 101 Market Street, San Francisco, California 94105-1579:

1. G V Bancorp, Inc., and G V Bancorp Employee Stock Ownership Plan, both of Gunnison, Utah; to become bank holding companies by acquiring 100 percent of the voting shares of Gunnison Valley Bank, Gunnison, Utah. Comments regarding this application must be received not later than March 2, 1998.

Board of Governors of the Federal Reserve System, February 3, 1998.

Jennifer J. Johnson,

Deputy Secretary of the Board.

[FR Doc. 98–3107 Filed 2–6–98; 8:45 am]

BILLING CODE 6210–01–F

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

The National Center for HIV, STD, and TB Prevention (NCHSTP): Meetings

Name: HIV Prevention Consultants Meetings.

Times and Dates: 9 a.m.-5 p.m., March 2, 1998. 9 a.m.-5 p.m., June 22, 1998. 9 a.m.-5 p.m., September 14, 1998. 9 a.m.-5 p.m., December 7, 1998.

Place: Washington Sheraton, 2660 Woodley Road, Washington, DC 20008, telephone 202/328–2000.

Status: Open to the public for observation and comment, limited only by the space available. The meeting rooms accommodate approximately 55 people.

Purpose: The purpose of these meetings is to provide a quarterly forum for consultations and discussion among representatives of governmental and nongovernmental organizations who are knowledgeable and experienced in HIV prevention policy, the staff of the Division of HIV/AIDS Prevention, and the National Center for HIV, STD, and TB Prevention.

Matters to be Discussed: Agenda items will include a discussion of broad HIV prevention programmatic and policy related issues.

Contact Person for More Information: Chad Martin, Division of HIV/AIDS Prevention, Intervention, Research, and Support NCHSTP, CDC, 1600 Clifton Road, M/S E–35, Atlanta, GA 30333, telephone 404/639–5200, email address: cgm8@cdc.gov

Dated: January 30, 1998.

Carolyn J, Russell,

Director, Management Analysis and Services Office, Centers for Disease Control and Prevention (CDC).

[FR Doc. 98–3154 Filed 2–6–98; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 98F–0058]

Sekisui Plastics Company, Ltd.; Filing of Food Additive Petition

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that Sekisui Plastics Co., Ltd., has filed a petition proposing that the food additive regulations be amended to expand the safe use of pyromellitic anhydride.

FOR FURTHER INFORMATION CONTACT: Mark A. Hepp, Center for Food Safety and Applied Nutrition (HFS–215), Food and Drug Administration, 200 C St. SW., Washington, DC 20204, 202–418–3098.

SUPPLEMENTARY INFORMATION: Under the Federal Food, Drug, and Cosmetic Act (sec. 409(b)(5)(21 U.S.C. 348(b)(5))), notice is given that a food additive petition (FAP 8B4582) has been filed by Sekisui Plastics Co., Ltd., c/o Bullwinkel Partners, Ltd., 19 S. LaSalle St., suite 1300, Chicago, IL 60603. The petition proposes to amend the food additive regulations in § 177.1630 *Polyethylene phthalate polymers* (21 CFR 177.1630) to expand the conditions of the safe use of pyromellitic anhydride as a modifier in ethylene terephthalate copolymers.

The agency has determined under 21 CFR 25.32(i) that this action is of a type that does not individually or cumulatively have a significant effect on the human environment. Therefore, neither an environmental assessment nor an environmental impact statement is required.

Dated: January 22, 1998.

Laura M. Tarantino,

Acting Director, Office of Premarket Approval, Center for Food Safety and Applied Nutrition.

[FR Doc. 98–3207 Filed 2–6–98; 8:45 am]

BILLING CODE 4160–01–F

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 98F–0054]

Sequa Chemicals, Inc.; Filing of Food Additive Petition

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that Sequa Chemicals, Inc., has filed a petition proposing that the food additive regulations be amended to provide for the safe use of octadecanoic acid, reaction products with 2-[(2-aminoethyl)amino]ethanol and urea, and the acetate salts thereof, which may be emulsified with ethoxylated tallow alkyl amines, for use in the manufacture of paper and paperboard intended for use in contact with dry food.

FOR FURTHER INFORMATION CONTACT: Vir D. Anand, Center for Food Safety and Applied Nutrition (HFS–216), Food and Drug Administration, 200 C St. SW., Washington, DC 20204, 202–418–3081.

SUPPLEMENTARY INFORMATION: Under the Federal Food, Drug, and Cosmetic Act (sec. 409(b)(5) (21 U.S.C. 348(b)(5))), notice is given that a food additive petition (FAP 8B4576) has been filed by Sequa Chemicals, Inc., c/o Keller and Heckman, 1001 G St. NW., suite 500 West, Washington, DC 20001. The petition proposes to amend the food additive regulations in § 176.180 *Components of paper and paperboard in contact with dry food* (21 CFR 176.180) to provide for the safe use of octadecanoic acid, reaction products with 2-[(2-aminoethyl)amino]ethanol and urea, and the acetate salts thereof, which may be emulsified with ethoxylated tallow alkyl amines, for increasing opacity and thickness, employed prior to the sheetforming operation in the manufacture of paper and paperboard intended for use in contact with dry food.

The agency has determined under 21 CFR 25.32(i) that this action is of a type that does not individually or cumulatively have a significant effect on the human environment. Therefore, neither an environmental assessment nor an environmental impact statement is required.